

THE CHRONICITY OF COGNITIVE IMPAIRMENT ASSOCIATED WITH EXPOSURE TO TOXIC MOLD

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INTRODUCTION

Gordon *et al.* (1999) reported an association between exposure to toxic molds and cognitive impairment. Since then, two other studies (Baldo, Ahmad, Ruff, 2002; Gordon, *et al.* 2004) have offered further support for such an association. While some efforts have been made to examine the chronicity of cognitive symptoms in samples of persons exposed to mold (e.g., Sudakin, 1998), there are no studies in the literature to date which examine patterns of performance over time on comprehensive batteries of neuropsychological tests. Examining chronicity is crucial to determining whether the cognitive impairments found to be associated with mold exposure persist for significant periods of time after exposure has ceased. The present study was conducted in order to begin to address this issue.

METHOD

A group of eight individuals with histories of exposure to toxic mold, who were originally administered a battery of neuropsychological tests by the senior author, were re-tested at an interval of one to five years after the initial testing. All eight were exposed to toxic molds such as *Stachybotrys atra*, *Penicillium*, and *Aspergillus* for varying lengths of time. All exposures were documented by environmental testing conducted during the exposure period. All exposures had ceased by the time of the first testing.

Seven of the eight persons tested were women. At the second testing, they ranged in age from 37 to 65 ($M = 49.25$, $SD = 10.12$). Most were college graduates (Mean years of education = 15.75, $SD = 1.95$). Participants' full scale IQs varied from the average to very superior ranges ($M = 116$, $SD = 11$). All eight continued to report symptoms of cognitive impairment at re-testing.

RESULTS

Because group data do not capture the full extent of the chronicity of cognitive impairments in a sample of this size, especially in higher functioning individuals, a brief description of each participant's functioning is provided below. In each case, although impairments were chronic in some domains, there was some fluctuation in test scores between evaluations. Some scores improved, some remained unimpaired, and some worsened. Because a thorough discussion of test and retest scores for each subject is beyond the scope of this paper, we have chosen to focus on domains where chronic impairments (either relative or absolute) were found at re-testing. Key supporting data for chronicity are provided in Table 1:

Participant 1 (Evaluated 1998 and 2002): Participant 1 is a professional with a master's degree. She continued to experience reduced processing speed, memory, and learning difficulties.

Participant 2 (Evaluated 1997 and 2001): Participant 2 graduated from college *cum laude*, with honors in two majors. Until the onset of her cognitive impairments, she worked in a literary field. At re-testing, verbal learning and memory continued to be reduced and verbal memory and IQ remained inconsistent with her levels of academic performance in college. Some difficulties with executive functions also remained at the second testing.

Participant 3 (Evaluated 2001 and 2002): Participant 3 continued to have significant impairments in the domains of attention and concentration and verbal learning at re-evaluation.

Participant 4 (Evaluated 2000 and 2002): On re-testing, Participant 4's test results continued to indicate reduced processing speed on cognitively demanding tasks.

Participant 5 (Evaluated 2001 and 2002): Participant 5's IQ is in the very superior range. Re-testing indicated the continuation of significant impairment on tests of memory function relative to IQ level. Difficulties on tests of executive functions also remained.

Participant 6 (Evaluated 2000 and 2002): At the second testing Participant 6 continued to have significant difficulties on tests of complex visual memory and impaired executive functions.

Participant 7 (Evaluated 1997 and 2002): Participant 7 is a college graduate who has a successful work history in the teaching, management, and financial fields. Re-test findings showed continued decreased intellectual functioning, impaired visual

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memory on complex tasks and reduced performance in aspects of executive functioning.

Participant 8 (Evaluated 1997 and 2002): Participant 8's IQ is in the superior range. On re-testing, relative impairments on tests of attention and concentration, visual memory, and verbal encoding were noted to have continued.

In all eight cases, a thorough review of relevant medical history revealed no preexisting problems that could account for the findings of cognitive impairment (e.g., traumatic brain injury, neurological disorders, significant psychiatric history) or for their chronicity.

DISCUSSION

These data indicate that, despite some variations in test findings, all participants continued to experience cognitive impairment one to five years after initial neuropsychological testing and termination of mold exposure. The findings suggest that some cognitive deficits secondary to toxic mold exposure are chronic in this small sample of individuals, even after cessation of mold exposure.

Table 1. Neuropsychological Testing Data^a from Study Participants Documenting Chronicity of Impairment from Initial Testing to Subsequent Re-testing

Participant	Percentile Scores on Testing 1	Percentile Scores on Testing 2
	Processing Speed	Processing Speed
1	WAIS-III ^b Processing Speed = 21%ile	WAIS-III Processing Speed = 21%ile
		Stroop ^c Word = 3%ile
		Stroop Color = 1%ile
		Trails A ^d = 6%ile
	Memory & Learning	Memory & Learning
	WAIS-III Working Memory = 9%ile	WAIS-III Working Memory = 18%ile
	WMS-III ^e Auditory Immediate = 13%ile	WMS-III Auditory Immediate = 9%ile
	WMS-III Auditory Delayed = 23%ile	WMS-III Auditory Delayed = 23%ile
	WMS-III Auditory Recognition = 25%ile	WMS-III Auditory Recognition = 37%ile
	WMS-III Visual Immediate = 7%ile	WMS-III Visual Immediate = 3%ile
2	WMS-III Visual Delayed = 50%ile	WMS-III Visual Delayed = 7%ile
	WMS-III Immediate memory = 5%ile	WMS-III Immediate memory = 3%ile
	WMS-III General memory = 27%ile	WMS-III General memory = 13%ile
	WMS-III Working memory = 5%ile	WMS-III Working memory = 8%ile
	CVLT ^f Trials 1 to 5 = <1%ile	CVLT Trials 1 to 5 = 2%ile
	Intellectual Functioning	Intellectual Functioning
	WAIS-III Verbal IQ = 82%ile	WAIS-III Verbal IQ = 77%ile
		WAIS-III VIQ < WAIS-III PIQ ($p < .05$)
	Verbal Learning & Memory	Verbal Learning & Memory
	WMS Verbal Memory Index = 97 (vs. Visual Memory Index = 135)	Auditory Immediate Memory = 63%ile
	CVLT Trial 5 = 2%ile	
	CVLT Trials 1 to 5 = 24%ile	
	All other CVLT scores = 16%ile	
	Executive Functions	Executive Functions
	Booklet category test = Could not complete	All CVLT scores = 50%ile
		Booklet category test = 8%ile

Percentile Scores on Testing 1	Percentile Scores on Testing 2
Attention & Concentration	Attention & Concentration
CPT ^g Overall Index = 10.67 (Borderline)	CPT Overall Index = 10.67 (Borderline)
CPT Confidence Index for	CPT Confidence Index for

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	Percentile Scores on Testing 1	Percentile Scores on Testing 2
3	Attention & Concentration CPT* Overall Index = 10.67 (Borderline) CPT Confidence Index for Attentional Problems = 93% Verbal Learning CVLT Trial 5 = 2%ile CVLT Long Delay Free Recall = 2%ile	Attention & Concentration CPT Overall Index = 10.67 (Borderline) CPT Confidence Index for Attentional Problems = 93% Verbal Learning CVLT Trial 5 = 6%ile CVLT Long Delay Free Recall = 6%ile
4	Processing Speed All PASAT* Scores < 1%ile WAIS-III Processing Speed Index = 21%ile Purdue pegboard scores = <2%ile	Processing Speed All PASAT Scores < 2%ile
5	Memory All WMS-III indexes except Working memory < WAIS-III VIQ ($p < .05$). All CVLT scores = 16 or 18%ile Executive Functions Booklet category Test = 42%ile Stroop Color/Word = 65%ile	Memory WMS-III Verbal Immediate and Delayed Memory < WAIS-III VIQ ($p < .05$). WMS-III Visual Immediate and Delayed Memory < WAIS-III PIQ ($p < .05$). Executive Functions Booklet category Test = 1%ile Trails B = 50%ile Stroop Color/Word = 58%ile
6	Visual Memory WMS-III Visual Immediate and Visual Delayed < than FSIQ ($p < .05$) CVLT Trials 1 to 5 < 16%ile Executive Functions Watson Glaser ¹ = 25%ile Booklet category test = 4%ile	Visual Memory All Rey Complex Figure scores = <1%ile Executive Functions Watson Glaser = 5%ile IOWA* reading comprehension = 30%ile (extra time required)

	Percentile Scores on Testing 1	Percentile Scores on Testing 2
7	Intellectual Functioning WAIS-R ¹ PIQ = 27%ile Visual Memory WMS-R ^m Visual Reproduction = 10%ile Executive Functions Watson Glaser = 3%ile Booklet category test = Could not complete	Intellectual Functioning WAIS-III PIQ = 34%ile Visual Memory Rey Complex Figure = <1%ile (Immediate & Delayed) Executive Functions IOWA reading comprehension = 16%ile IOWA reading efficiency = 6%ile Watson Glaser = 10%ile Stroop Color Word = 24%ile
8	Attention & Concentration WMS-R Attention/ Concentration Index Score = 89 Visual Memory WMS-R Visual Memory Index = 113 Verbal Encoding CVLT Trial 1 = 50%ile	Attention & Concentration CPT Clinical Significant Attention Problem, Confidence Index = 99.9% Visual Memory WMS-III Visual Immediate Memory < WMS-III Auditory and Visual Delayed Memory (p < .05) Verbal Encoding CVLT Trial 1 = 50%ile

*Only data indicative of significant impairments (absolute or relative) where performance in the given domain did not change significantly from Testing 1 to Testing 2 are included, ^b Wechsler Adult Intelligence Scale - Third Edition, ^c Wechsler Memory Scale - Third Edition, ^d California Verbal Learning Test, ^e Stroop Color Word Test, ^f Trail Making Test - Part A, ^g Conners Continuous Performance Test, ^h Paced Auditory Serial Attention Task, ⁱ Trail Making Test - Part B, ^j Watson Glaser Critical Thinking Appraisal - Form B, ^k Iowa Silent Reading Test, ^l Wechsler Adult Intelligence Scale - Revised, ^m Wechsler Memory Scale - Revised

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